

PRE-APPEAL BRIEF REQUEST FOR REVIEW		Docket Number (Optional) 133172.02201							
I hereby certify that this correspondence is being deposited with the United States Postal Service with sufficient postage as first class mail in an envelope addressed to "Mail Stop AF, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450" [37 CFR 1.8(a)] on _____ Signature _____ Typed or printed name _____		<table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%; padding: 5px;"> Application Number 10518701 </td> <td style="width: 50%; padding: 5px;"> Filed September 1, 2005 </td> </tr> <tr> <td colspan="2" style="padding: 5px;"> First Named Inventor Arnold I. Levinson </td> </tr> <tr> <td style="padding: 5px;"> Art Unit 1644 </td> <td style="padding: 5px;"> Examiner Michael E. Szperka </td> </tr> </table>		Application Number 10518701	Filed September 1, 2005	First Named Inventor Arnold I. Levinson		Art Unit 1644	Examiner Michael E. Szperka
Application Number 10518701	Filed September 1, 2005								
First Named Inventor Arnold I. Levinson									
Art Unit 1644	Examiner Michael E. Szperka								
<p>Applicant requests review of the final rejection in the above-identified application. No amendments are being filed with this request.</p> <p>This request is being filed with a notice of appeal.</p> <p>The review is requested for the reason(s) stated on the attached sheet(s). Note: No more than five (5) pages may be provided.</p>									
I am the <input type="checkbox"/> applicant/inventor. <input type="checkbox"/> assignee of record of the entire interest. See 37 CFR 3.71. Statement under 37 CFR 3.73(b) is enclosed. (Form PTO/SB/96) <input checked="" type="checkbox"/> attorney or agent of record. Registration number <u>52,201</u> <input type="checkbox"/> attorney or agent acting under 37 CFR 1.34. Registration number if acting under 37 CFR 1.34 _____		<table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%; padding: 5px;"> /Daniel M. Scolnick, Reg. No. 52,201/ Signature Daniel M. Scolnick Typed or printed name 610-640-7820 Telephone number July 23, 2009 Date </td> <td style="width: 50%;"></td> </tr> </table>		/Daniel M. Scolnick, Reg. No. 52,201/ Signature Daniel M. Scolnick Typed or printed name 610-640-7820 Telephone number July 23, 2009 Date					
/Daniel M. Scolnick, Reg. No. 52,201/ Signature Daniel M. Scolnick Typed or printed name 610-640-7820 Telephone number July 23, 2009 Date									
NOTE: Signatures of all the inventors or assignees of record of the entire interest or their representative(s) are required. Submit multiple forms if more than one signature is required, see below*.									
<input type="checkbox"/> *Total of _____ forms are submitted.									

This collection of information is required by 35 U.S.C. 132. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11, 1.14 and 41.6. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Mail Stop AF, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

If you need assistance in completing the form, call 1-800-PTO-9199 and select option 2.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patent application or patent. Accordingly, pursuant to the requirements of the Act, please be advised that: (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) furnishing of the information solicited is voluntary; and (3) the principal purpose for which the information is used by the U.S. Patent and Trademark Office is to process and/or examine your submission related to a patent application or patent. If you do not furnish the requested information, the U.S. Patent and Trademark Office may not be able to process and/or examine your submission, which may result in termination of proceedings or abandonment of the application or expiration of the patent.

The information provided by you in this form will be subject to the following routine uses:

1. The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether disclosure of these records is required by the Freedom of Information Act.
2. A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiations.
3. A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
4. A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552a(m).
5. A record related to an International Application filed under the Patent Cooperation Treaty in this system of records may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant to the Patent Cooperation Treaty.
6. A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or his/her designee, during an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2904 and 2906. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
8. A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S.C. 122(b) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspection or an issued patent.
9. A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of: Arnold I. Levinson *et al.*

Serial No.: 10/518,701

Filed: September 1, 2005

Group Art Unit: 1644

Examiner: Michael E. Szperka

Confirmation No.: 5645

Title: VACCINES FOR SUPPRESSING IGE-MEDIATED ALLERGIC DISEASE AND METHODS FOR USING THE SAME

Mail Stop: AF

Commissioner for Patents

P.O. Box 1450

Alexandria, VA 22313-1450

Pre-Appeal Brief Request for Review

Dear Sir:

In response to the Final Rejection dated January 23, 2009 and the Advisory Action dated April 7, 2009, Applicants respectfully request reconsideration of the pending rejections.

I. Claims 1-3, 5-7, 22-24, and 26-29 stand rejected under 35 U.S.C. § 103(a) as allegedly being unpatentable over Chen *et al.* (WO 98/53843) in view of Wang *et al.* (WO 99/67293) in view of Hollis *et al.* (U.S. Patent No. 5,629,415) and in view of Rutter (U.S. Patent No. 4,769,326). The Office alleges that the Chen reference discloses vaccine constructs comprising the membrane bound domain of IgE coupled to heterologous sequences including helper T epitopes. The Office acknowledges that the Chen reference fails to disclose a composition with a proteolytic cleavage sequence. The Office alleges the deficiency is remedied by the Rutter reference which allegedly discloses the use of linkers comprising proteolytic cleavage sites because the linkers “allow for efficient incorporation and removal of desired functional properties.” (Final Office Action p. 5). Applicants respectfully disagree.

The office has clearly erred by not properly establishing a *prima facie* case of obviousness and because the cited references teach away from the claimed invention. The Chen and Wang references teach away from using a proteolytic cleavage sequence, which would allow the components of the construct to be unconjugated. One of skill in the art reading the Chen and Wang references would be led to use a composition that conjugates the two components without the two components being able to be cleaved. For example, the Chen reference repeatedly describes conjugates and does not state that the conjugated composition can include a cleavage

sequence. Additionally, the Chen reference states that for conjugates in human use one would expect that there would be “no inhibition of IgE responses to unrelated, *unconjugated* antigens.” (Chen, p. 10, line 22, emphasis added.). ““A reference may be said to teach away when a person of ordinary skill, upon reading the reference, would be discouraged from following the path set out in the reference, or would be led in a direction divergent from the path that was taken by the applicant.”” *Optivus Tech., Inc. v. Ion Beam Applications S.A.*, 469 F.3d 978, 989 (Fed. Cir. 2006) (quoting *In re Kahn*, 441 F.3d 977, 990 (Fed. Cir. 2006)). Here, the Chen reference teaches away because it states that there would be no inhibition of IgE responses for unconjugated antigens. Therefore, one of skill in the art reading the Chen reference in its entirety would not have inserted a proteolytic cleavage sequence because such a construct would lead to an unconjugated composition leading to a result that is not desired. Accordingly, the Chen reference teaches away from using a construct that would allow the epitopes to be separated by use of a cleavage sequence.

The Wang reference also teaches away from including a proteolytic cleavage sequence. The Wang reference discloses the use of a spacer between its components and states that the two components are “adjacent to either the N- or C-terminus of IgE-CH3 domain antigen sequences, *in order to evoke efficient* antibody responses.” (Wang, p. 28-29). Like the Chen reference, the Wang reference teaches that the components should be next to one another and linked so that there is a proper response. The Wang reference fails to suggest decoupling the components and teaches away from such a method because it would not “evoke [an] efficient antibody response[.]”

The Advisory Action argues that the proteolytic cleavage sequence may not be cleaved *in vivo* and that separation is not a limitation of the claim. The Office’s assertion is reading the limitation out of the claim. The cleavage sequence is a functional cleavage sequence otherwise the term cleavage sequence would have no meaning. One of skill in the art reading the references would not have added a cleavage sequence because the purpose of including a cleavage sequence is to allow the parts to be separated, which is exactly what the references teach away from. Accordingly, one of skill in the art would not have used a proteolytic cleavage sequence because the Wang and Chen references teach away from allowing the components to be separated. In view of the foregoing, Applicants respectfully request that the rejection under 35 U.S.C. § 103(a) be withdrawn.

II. Claims 1-3, 5-7, 22-24, and 26-29 stand rejected under 35 U.S.C. § 103(a) as allegedly being unpatentable over Klysner *et al.* (WO 02/20038) in view of Wang *et al.* (WO 99/67293) and in view of Rutter. Claims 1-3, 5-7, 22-24, and 26-29 stand rejected under 35 U.S.C. § 103(a) as allegedly being unpatentable over Klysner *et al.* (US 2002/0172673) in view of Wang *et al.* (WO 99/67293) and in view of Rutter. The Office acknowledges that the Klysner reference fails to disclose the use of linkers comprising a proteolytic cleavage sequence between the epitopes. (Office Action, page 6). The Office alleges that the Rutter reference cures this deficiency. Applicants respectfully disagree.

The Office has clearly erred because the claims are not obvious because the Klysner and Wang references each teach away from inserting a proteolytic cleavage sequence. The Wang reference teaches away for the reasons stated above. The Klysner reference also teaches away because the Klysner reference teaches that the epitopes described in Klysner should be simultaneously presented by the antigen presenting cells. (Klysner, p. 13, lines 14-20)¹. The inclusion of a proteolytic cleavage sequence that allows the epitopes to be separated would function to reduce the likelihood of simultaneous presentation of the epitopes by the antigen presenting cells. One of skill in the art would not have been led to insert a proteolytic cleavage sequence because it would be contrary to what the Klysner reference states is necessary for an effective use, that is the simultaneous presentation of the epitopes by the antigen presenting cells. Thus, the Klysner reference teaches away from using a proteolytic cleavage sequence.

The Advisory Action argues that the proteolytic cleavage sequence may not be cleaved *in vivo* and that separation is not a limitation of the claim. The Office's assertion is reading the limitation out of the claim. The cleavage sequence is a functional cleavage sequence otherwise the term cleavage sequence would have no meaning. One of skill in the art reading the references would not have added a cleavage sequence because the purpose of including a cleavage sequence is to allow the parts to be separated, which is exactly what the references teach away from. Accordingly, one of skill in the art would not have used a proteolytic cleavage sequence because the Klysner and Wang references teach away from allowing the components to be separated, and, therefore, the references teach away from including a cleavage sequence.

¹ Applicants note that the two Klysner references are the same. Page and line number refer to WO 02/20038

Therefore, the claims are not obvious because the Klysner and Wang references teach away and because the combination of the references would not have suggested one of skill in the art to insert a proteolytic cleavage sequence. In view of the foregoing, Applicants respectfully request that the rejection under 35 U.S.C. § 103(a) be withdrawn.

III. Claims 8, 32-37, 50, 58-73 stand rejected under 35 U.S.C. § 103(a) as allegedly being unpatentable over Chen et al (WO 98/53843) in view of Wang et al. (WO 96/67293) in view of Hollis et al. (U.S. Patent No. 5,629,415) and in view of Rutter (U.S. Patent 4,769,326) as applied to claims 1-3, 5-7, 22-24, and 26-29, and further in view of Walls et al. (Nucleic Acids Research, 1993, 21:2921-2929) as evidenced by Janeway et al (Immunobiology, 3rd Edition, Garland Publications, 1997, pages 3:26-3:31). Claims 8, 32-37, 50, 58-73 stand rejected under 35 U.S.C. § 103(a) as allegedly being unpatentable over Klysner (WO 02/20038) in view of Wang et al. (WO 96/67293) in view of Hollis et al. (U.S. Patent No. 5,629,415) and in view of Rutter (U.S. Patent 4,769,326) as applied to claims 1-3, 5-7, 22-24, and 26-29, and further in view of Walls et al. (Nucleic Acids Research, 1993, 21:2921-2929) as evidenced by Janeway et al (Immunobiology, 3rd Edition, Garland Publications, 1997, pages 3:26-3:31). Claims 8, 32-37, 50, 58-73 stand rejected under 35 U.S.C. § 103(a) as allegedly being unpatentable over Klysner (US 2002/0172673) in view of Wang et al. (WO 96/67293) in view of Hollis et al. (U.S. Patent No. 5,629,415) and in view of Rutter (U.S. Patent 4,769,326) as applied to claims 1-3, 5-7, 22-24, and 26-29, and further in view of Walls et al. (Nucleic Acids Research, 1993, 21:2921-2929) as evidenced by Janeway et al (Immunobiology, 3rd Edition, Garland Publications, 1997, pages 3:26-3:31). Applicants respectfully disagree and assert that the Office has clearly erred because the combination fails to yield the claimed invention.

The Office alleges that the combination of references discloses each and every element of claims 8, 32-37, 50, 58-73. In support of this contention the Office states:

Note that as evidenced by Janeway et al., immunoglobulin genes are assembled via the process of V(DJ) recombination, and that different isotypes (i.e. IgG, IgE, IgA) are obtained by isotype switching. As such the immunoglobulin heavy chain leader sequence is upstream of the rearranged variable domain . . . and thus an "IgE leader" is the *same sequence* as an IgM, IgD, IgG, and IgA leader sequence...Thus, the "Ig leader" of Walls *et al.* is an "IgE leader."

(Final Office Action, pages 9-10, emphasis added). Applicants previously submitted a declaration pursuant to 37 C.F.R. § 1.132 by Dr. David B. Weiner. The declaration lists the amino acid leader sequences from sequences that have been labeled as IgE variable, IgA constant, IgA, variable 1, IgA variable 2, IgA variable 3, IgG constant, IgM variable, and IgM VH1. The declaration also shows the sequence similarity between the different leader sequences. The declaration shows that the IgE leader sequence is not 100% identical to the other leader sequences. The declaration states, “[t]he alignments show that IgE leader sequence is not the same as the leader sequences from the different isotypes.” (Declaration, ¶ 3). Therefore, the “Ig leader” of the Walls reference is not an “IgE leader.” Therefore, the Office has failed to present a proper *prima facie* obviousness rejection because even if all the references were combined the combination does not yield the present invention.

The Office has rejected the Declaration because of the apparent lack of Genbank or other source identifiers. The Declaration must be treated as a fact and as such Applicant has provided evidence showing that not all leader sequences are the same. The Office has not provided any evidence to show that the facts presented in the declaration are incorrect. The sequences demonstrate that not all leader sequences are the same, and, therefore, the rejection is improper for the reasons stated above.

In view of the foregoing, Applicants respectfully request that the rejection under 35 U.S.C. § 103(a) be withdrawn.

Applicants respectfully submit that the claims are in condition for allowance. The Office is invited to contact Applicants’ undersigned representative at 610-640-7820 to resolve any remaining issues. The Commissioner is hereby authorized to charge any deficiencies of fees and credit of any overpayments to Deposit Account No. 50-0436.

Respectfully Submitted,
/Daniel M. Scolnick, Reg. No. 52,201/
Daniel M. Scolnick, Ph.D.
Registration No. 52,201

Dated: **July 23, 2009**
PEPPER HAMILTON, LLP
400 Berwyn Park
899 Cassatt Road
Berwyn, PA 19312-1183
Telephone: 610-640-7820
Facsimile: 610-640-7835